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detecting electrode 67 and the second working electrode 69 are connected with a bias voltage 71.

A particular working process is as follows. On the surface of the CEA-LAPS sensor 68 are disposed the nanolayer 75 and the biotin layer 76. At this point, the condensate flows into the CEA detecting cavity, the CEA antigen 77 in the condensate bonds with the biotin layer 76 via a covalent bond, and affixes to the surface of the CEA-LAPS sensor 68. Then, the CEA antibody-avidin 79 and the CEA antibody-urease 78 flow into the CEA detecting cavity, and bonded with the CEA antigen 77, forming a sandwich structure as shown in the figure. Then, the washing liquid cleans out extra CEA antibody-urease 78, and only leaves the sandwich structure consisting of the CEA antigen 77, the CEA antibody-urease 78 and the CEA antibody-avidin 79 on the CEA-LAPS sensor 68 in the CEA detecting cavity. Then, the urea flows into the CEA detecting cavity, reacts with the urease on the sandwich structure, and leads to a change of pH, and the change in pH is related with the amount of the CEA antigen 77 in the condensate. The CEA-LAPS sensor 68 obtains the amount of the CEA antigen 77 in the condensate by detecting the change of pH.

The invention claimed is:

1. An integrated analysis device for simultaneously detecting human exhaled breath condensates (EBCs) and volatile organic compounds (VOCs) in human exhaled breath, comprising a module for sampling, separating and enriching a detected object, an EBCs detection module and a combined VOCs detection module; the module for sampling, separating and enriching the detected object is connected with the EBCs detection module via a syringe pump for sample injection, and the module for sampling, separating and enriching the detected object is connected with the combined VOCs detection module via a capillary separation column; wherein the EBCs detection module comprises an EBC inlet (57), an inlet for washing liquid (58), a first three-way valve (59), a composite light addressable potentiometric sensor (LAPS) for heavy metal ions ( $\text{Cr}^{3+}$ ) (60), a first working electrode (61), a light source controlled by a signal generating circuit (62), a reference electrode (63), a second three-way valve (64), a urea inlet (65), a detecting electrode (67), a carcinoembryonic antigen light addressable potentiometric sensor (CEA-LAPS) (68), a second working electrode (69), a  $\text{Cr}^{3+}$  ion detecting cavity and a carcinoembryonic antigen (CEA) detecting cavity; the EBC inlet (57), the inlet for washing liquid (58) and the  $\text{Cr}^{3+}$  ion detecting cavity are connected via the first three-way valve (59); the urea inlet (65), the  $\text{Cr}^{3+}$  ion detecting cavity and the CEA detecting cavity are connected via the second three-way valve (64); the reference electrode (63) is inserted into the  $\text{Cr}^{3+}$  ion detecting cavity from its top; the composite LAPS for heavy metal ions ( $\text{Cr}^{3+}$ ) (60) and the first working electrode (61) are fixed to the bottom of the  $\text{Cr}^{3+}$  ion detecting cavity; the first working electrode (61) is joined with the bottom of the composite LAPS for heavy metal ions ( $\text{Cr}^{3+}$ ) (60); the detecting electrode (67) is inserted into the CEA detecting cavity from its top; the CEA-LAPS (68) and the second working electrode (69) are fixed to the bottom of the CEA detecting cavity, and the second working electrode (69) is joined with the bottom of the CEA-LAPS (68); at the upper portion of the CEA detecting cavity are disposed an inlet for CEA antibody-urease compound liquid (66) and an outlet for waste liquid (70); one light source controlled by the signal generating circuit (62) is placed at a position corresponding to the composite LAPS for heavy metal ions ( $\text{Cr}^{3+}$ ) (60) under the  $\text{Cr}^{3+}$  ion detecting cavity, and another light source controlled by the signal generating circuit (62) is

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placed at a position corresponding to the CEA-LAPS (68) under the CEA detecting cavity.

2. The integrated analysis device for simultaneously detecting EBCs and VOCs in human exhaled breath according to claim 1, wherein the CEA-LAPS (68) is built by depositing a silicon dioxide ( $\text{SiO}_2$ ) layer (73) and a silicon nitride ( $\text{Si}_3\text{N}_4$ ) film (74) in turn on a Si substrate (72) a chemical vapor deposition and photolithography, and forming a nanolayer (75) and a biotin layer (76) on the surface of the  $\text{Si}_3\text{N}_4$  film (74) by a chemical coating method.

3. The integrated analysis device for simultaneously detecting EBCs and VOCs in human exhaled breath according to claim 1, the module for sampling, separating and enriching the detected object further comprising a mouthpiece (1), a saliva collector (2), an inspiration check valve (3), an expiration check valve (4), a first three-way solenoid valve (5), a second three-way solenoid valve (6), an activated carbon filter (7), an inlet check valve (8), a gas buffering chamber (9), a third three-way solenoid valve (10), a gas outlet (11), an adsorption tube (12), a miniature vacuum pump (13), a gas mass flow meter (14), a linear stepping motor (15), a piston (16), a washing liquid storage for condensation tube (17), a peristaltic pump (18), a condensation module (19), and a condensate collector (20); wherein the mouthpiece (1) is connected with the first three-way solenoid valve (5) via the inspiration check valve (3) and the expiration check valve (4) in turn, the saliva collector (2) is connected between the mouthpiece (1) and the inspiration check valve (3), the two outlets of the first three-way solenoid valve (5) are connected respectively with the second three-way solenoid valve (6) and the gas mass flow meter (14); the inlet check valve (8) is then connected with the third three-way solenoid valve (10) via the activated carbon filter (7), the second three-way solenoid valve (6) and the gas buffering chamber (9) in turn, the two outlets of the third three-way solenoid valve (10) is then connected respectively with the gas outlet (11) and the adsorption tube (12) and to the miniature vacuum pump (13); the gas mass flow meter (14) is connected with the condensation module (19), and all the above connections are a gas pipeline connection; the washing liquid storage for condensation tube (17) is connected with the condensate collector (20) in a liquid pipeline via the peristaltic pump (18) and the condensation module (19) in turn; the linear stepping motor (15) is connected with the piston (16), and drives the piston (16) into the condensation module (19).

4. The integrated analysis device for simultaneously detecting EBCs and VOCs in human exhaled breath according to claim 3, the gas buffering chamber (9) further comprising a heating rod (21), a heating piece (22) and a conduit (23), wherein both the heating rod (21) and the conduit (23) are placed within the heating piece (22).

5. The integrated analysis device for simultaneously detecting EBCs and VOCs in human exhaled breath according to claim 3, the condensation module (19) further comprising the piston (16), a gas inlet (26), an inlet for washing liquid (27), a condensation tube (28), an ice cooling box (29), and a condensate outlet (30); the gas inlet (26), the inlet for washing liquid (27), the condensate outlet (30) and a condensation tube (28) are formed integrally, the piston (16) is inserted into the condensation tube (28), and the ice cooling box (29) is in close contact with the outside of the condensation tube (28).

6. The integrated analysis device for simultaneously detecting EBCs and VOCs in human exhaled breath according to claim 3, the module for sampling, separating and enriching the detected object further comprising an aluminium piece (24) and a cooling plate (25), the aluminium piece (24) is in close contact with the periphery of the adsorp-